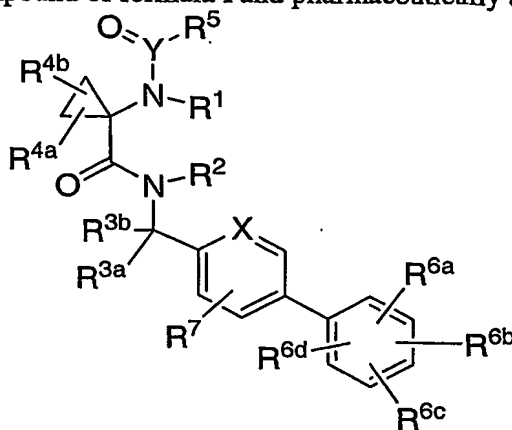


WHAT IS CLAIMED IS:

1. A compound of formula I and pharmaceutically acceptable salts thereof:



I

wherein

R¹ and R² are independently selected from hydrogen and C₁₋₄ alkyl;

R^{3a} and R^{3b} are independently selected from hydrogen and C₁₋₄ alkyl optionally substituted with 1 to 5 halogen atoms;

- 10 R^{4a} and R^{4b} are independently selected from hydrogen, halogen, and C₁₋₄ alkyl optionally substituted with 1 to 4 groups selected from halogen, OR^a, OC(O)R^a, S(O)_kR^d, OS(O)₂R^d, and NR¹R², or R^{4a} and R^{4b} together with the carbon atom to which they are both attached form an exo-cyclic methylene optionally substituted with 1 to 2 groups selected from C₁₋₄ alkyl optionally substituted with 1-5 halogens and C₁₋₄ alkyloxy;

- 15 R⁵ is selected from (1) C₁₋₆ alkyl optionally substituted with 1 to 5 groups independently selected from halogen, nitro, cyano, OR^a, SR^a, COR^a, SO₂R^d, CO₂R^a, OC(O)R^a, NR^bR^c, NR^bC(O)R^a, NR^bC(O)₂R^a, C(O)NR^bR^c, C₃₋₈ cycloalkyl, (2) C₃₋₈ cycloalkyl optionally substituted with 1 to 5 groups independently selected from halogen, nitro, cyano and phenyl, (3) C₃₋₆ alkynyl, (4) C₂₋₆ alkenyl optionally substituted with hydroxyethyl, (5) (CH₂)_k-aryl optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, OR^a, SR^a, C(O)₂R^a, C₁₋₄ alkyl and C₁₋₃ haloalkyl; 20 (6) (CH₂)_k-heterocycle optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, OR^a, SR^a, C₁₋₄ alkyl and C₁₋₃ haloalkyl wherein said heterocycle is selected from (a) a 5-membered heteroaromatic ring having a ring heteroatom selected from N, O and S, and optionally having up to 3 additional ring nitrogen atoms wherein said ring is optionally benzo-fused; (b) a 6-membered heteroaromatic ring containing from 1 to 3 ring nitrogen atoms and N-oxides thereof, wherein said ring is 25 optionally benzo-fused; and (c) a 5- or 6-membered non-aromatic heterocyclic ring selected from

tetrahydrofuranyl, 5-oxotetrahydrofuranyl, 2-oxo-2H-pyranyl, 6-oxo-1,6-dihydropyridazinyl, (7) $C(O)_2R^a$, and (8) $C(O)NR^bR^c$;

R^{6a} is selected from (1) $-OSO_2R^8$, (2) $-NR^{8a}SO_2R^9$, and (3) $-C(R^{8b})(R^{8c})SO_2R^9$;

R^{6b} , R^{6c} , and R^{6d} are independently selected from (1) hydrogen, (2) halogen, (3) OSO_2R^8 , (4) C_{1-4} alkyl optionally substituted with 1 to 5 halogen atoms, (5) cyano, (6) nitro, (7) OR^a , and (8) CO_2R^a , or when attached to adjacent carbon atoms R^{6c} and R^{6d} together with the carbon atoms to which they are attached form a 5- to 8-membered saturated or unsaturated ring;

R^7 is selected from (1) hydrogen, (2) halogen, (3) cyano, (4) nitro, (5) OR^a , (6) CO_2R^a , (7) $C(O)NR^bR^c$, and (8) C_{1-4} alkyl optionally substituted with 1 to 5 halogen atoms,

R^8 is selected from (1) C_{1-4} alkyl optionally substituted with 1 to 5 halogen atoms, (2) $(CH_2)_k$ -aryl optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, $NR^aC(O)R^a$, OR^a , SR^a , CO_2R^a , C_{1-4} alkyl, C_{1-3} haloalkyl and NR^bR^c , (3) NR^bR^c , and (4) hydrogen;

R^{8a} is selected from hydrogen, C_{1-4} alkyl optionally substituted with 1 to 5 halogen atoms, halogen, and CO_2R^a , or

when R^{6a} and R^{6b} are attached to adjacent atoms, R^{8a} and R^{6b} together complete 5- or 6-membered ring;

R^{8b} and R^{8c} are independently selected from hydrogen, C_{1-4} alkyl optionally substituted with 1 to 5 halogen atoms, halogen, cyano, nitro, CO_2R^a , and OR^a ;

R^9 is selected from (1) C_{1-4} alkyl optionally substituted with 1 to 5 halogen atoms, (2) aryl optionally

substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, $NR^aC(O)R^a$, OR^a , SR^a , CO_2R^a , C_{1-4} alkyl and C_{1-3} haloalkyl, and (3) $(CH_2)_k$ -aryl optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, $NR^aC(O)R^a$, OR^a , SR^a , $C(O)_2R^a$, C_{1-4} alkyl and C_{1-3} haloalkyl, or

R^{8a} and R^9 together with the atoms to which they are attached form a 5- to 8-membered heterocyclic ring;

R^a , R^b and R^c are independently selected from (1) hydrogen, (2) C_{1-4} alkyl optionally substituted with 1 to 5 halogen atoms, (3) phenyl optionally substituted with 1 to 3 groups selected from halogen, cyano, nitro, OH, C_{1-4} alkyloxy, C_{3-6} cycloalkyl and C_{1-4} alkyl optionally substituted with 1 to 5 halogen atoms, and (4) C_{3-6} cycloalkyl, or

R^b and R^c together with the nitrogen atom to which they are attached form a 4-, 5-, or 6-membered ring optionally containing an additional heteroatom selected from N, O, and S; or

R^b and R^c together with the nitrogen atom to which they are attached form a cyclic imide;

R^d is selected from (1) C_{1-4} alkyl optionally substituted with 1 to 5 halogen atoms, (2) C_{1-4} alkyloxy, (3) phenyl optionally substituted with 1 to 3 groups selected from halogen, cyano, nitro, OH, C_{1-4}

alkyloxy, C₃₋₆ cycloalkyl and C₁₋₄ alkyl optionally substituted with 1 to 5 halogen atoms, and (4) hydrogen;

X is selected from CH and N;

Y is selected from C and S=O; and

5 k is selected from 0, 1, and 2.

2. A compound of Claim 1 wherein R⁵ is selected from pyrimidinyl and C₁₋₆ alkyl optionally substituted with 1 to 5 groups independently selected from halogen.

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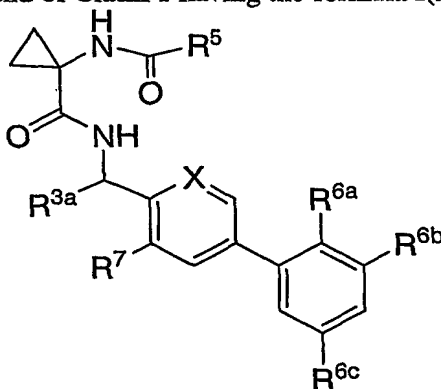
3. A compound of Claim 1 wherein Y is C.

4. A compound of Claim 1 wherein R^{6a} is OSO₂R⁸ and R⁸ is selected from 2,2,2-trifluoroethyl, trifluoromethyl, methyl, ethyl, propyl, isopropyl, phenyl, benzyl, and dimethylamino; or
15 R^{6a} is NHSO₂R⁹ and R⁹ is methyl or trifluoromethyl.

5. A compound of Claim 1 wherein R^{6b} is selected from hydrogen, fluorine, and chlorine.

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6. A compound of Claim 1 having the formula I(2):



I(2)

wherein X is N or CH, R^{3a} is H or C₁₋₄alkyl, R⁷ is hydrogen or halogen, and R⁵, R^{6a}, R^{6b} and R^{6c} have the same definitions as provided in Claim 1.

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7. A compound of Claim 6 wherein R^{6a} is NHSO₂R⁹; R⁹ is C₁₋₄alkyl optionally substituted with 1 to 5 halogen atoms, R^{6b} is halogen, and R^{6c} is hydrogen or halogen.

8. A compound of Claim 10 wherein R^{6a} is OSO₂R⁸; R⁸ is selected from methyl, trifluoromethyl, ethyl, propyl, isopropyl, benzyl, dimethylamino, 2,2,2-trifluoroethyl, and phenyl; R^{6b} is hydrogen or halogen, and R^{6c} is hydrogen or halogen.

9. A compound of Claim 10 wherein R⁵ is pyrimidinyl or C₁₋₄alkyl optionally substituted with 1 to 5 groups independently selected from halogen.

10. A compound selected from
 3,3'-difluoro-4'-{[(1-[(pyrimidin-5-ylcarbonyl)amino]cyclopropyl)carbonyl]amino}methyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoromethyl)sulfonyl]amino}cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,
 1-[(1R)-1-(3,3'-difluoro-2'-{[(trifluoromethyl)sulfonyl]oxy}-1,1'-biphenyl-4-yl)ethyl]amino}carbonyl)-cyclopropanaminium trifluoroacetate,
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl methanesulfonate,
 5-chloro-3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl ethanesulfonate,
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl propane-1-sulfonate
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl propane-2-sulfonate,
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl benzenesulfonate,
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl phenylmethanesulfonate

3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl)amino]ethyl}-1,1'-biphenyl-2-yl dimethylsulfamate,

3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl)amino]ethyl}-1,1'-biphenyl-2-yl 2,2,2-trifluoroethanesulfonate,

5 3-chloro-3'-fluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl)amino]ethyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,

3'-fluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl)amino]ethyl}-2-[(trifluoromethyl)sulfonyl]oxy}-1,1'-biphenyl-3-yl trifluoromethanesulfonate,

10 N-(1-[(1-[(1R)-1-(3,3'-difluoro-2'-[methyl(methylsulfonyl)amino]-1,1'-biphenyl-4-yl)ethyl)amino]carbonyl]cyclopropyl)pyrimidine-5-carboxamide,

N-(1-[(1-[(3,3'-difluoro-2'-[(methylsulfonyl)amino]-1,1'-biphenyl-4-yl)methyl)amino]carbonyl]cyclopropyl)pyrimidine-5-carboxamide,

N-{1-[(1-[(2'-(1,1-dioxido-1,2-thiazinan-2-yl)-3,3'-difluoro-1,1'-biphenyl-4-yl)methyl]amino)carbonyl]cyclopropyl}pyrimidine-5-carboxamide,

15 N-[(1R)-1-(3,3'-difluoro-2'-[(trifluoromethyl)sulfonyl]methyl)-1,1'-biphenyl-4-yl)ethyl]-1-[(trifluoroacetyl)amino]cyclopropanecarboxamide,

N-[(1R)-1-(3,3'-difluoro-2'-[(trifluoromethyl)sulfonyl]amino)-1,1'-biphenyl-4-yl)ethyl]-1-[(trifluoroacetyl)amino]cyclopropanecarboxamide, and

20 N-(1-[(1-[(1R)-1-(3,3'-difluoro-2'-[(methylsulfonyl)amino]-1,1'-biphenyl-4-yl)ethyl)amino]carbonyl]cyclopropyl)pyrimidine-5-carboxamide.

11. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 1 and a pharmaceutically acceptable carrier.

25 12. Use of a compound of Claim 1 or a pharmaceutically acceptable salt thereof in the manufacture of a medicament for the treatment or prevention of pain and inflammation.

13. Use of Claim 30 wherein said pain is postherpetic neuropathy, osteoarthritis pain, or dental pain.

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